

# REPORT DOCUMENTATION PAGE

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*CJ*

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✓ Spreadsheet  
✓ DTS

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FROM: PROI (TI) (STINFO)

28 May 1999

SUBJECT: Authorization for Release of Technical Information, Control Number: AFRL-PR-ED-TP-FY99-0115  
DeRose and Fajardo, "HEDM Source Characterization by Multi-Photon Ionization Time-of-Flight Mass Spectrometry"

Presentation **HEDM Conference**

(Statement A)

HEDM Source Characterization by  
Multi-Photon Ionization  
Time-Of-Flight Mass Spectrometry

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## Abstract

Thermochemical calculations show that a High Energy Density Matter (HEDM) fuel consisting of 5% boron atoms in cryogenic solid hydrogen would provide a 21% (or 80 seconds) improvement in Isp over LOX/LH<sub>2</sub>.<sup>1</sup> However, to date only about 0.1% metal atom concentration has been achieved in a solid hydrogen matrix. These studies have demonstrated that low concentrations of HEDM species are chemically stable in solid hydrogen. However, in order to obtain higher HEDM species concentrations we must determine optimum matrix deposition conditions. Knowing the dopant species' identities before and after deposition will quantify HEDM species recombination during the deposition process, and will facilitate the desired optimization.

We present results obtained using an apparatus designed to characterize the species produced by a variety of HEDM sources. In this apparatus, the HEDM species are ionized by a pulsed excimer laser beam and analyzed by time-of-flight mass spectrometry. Complications arising from photofragmentation vs. photoionization were encountered and documented as a function of ionization wavelength and intensity.

Data from experiments with HEDM precursor-coated tungsten filament sources are presented. These sources have numerous advantages, including low cost and versatility. Boron atom sources are made by chemical vapor deposition of pyrolyzed  $B_2H_6$  onto a filament. Characterization of the gas phase products of these sources indicates significant contamination by  $B_xH_y$  species.

The time-of-flight apparatus will be used to assist in the development and to verify the operation of future HEDM sources. By comparing what is produced by a source with what is trapped in the matrix, we can obtain a better understanding of the deposition process and insight into obtaining the desired higher concentration HEDM matrices.

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1. Carrick, P.G., "Specific Impulse Calculations of High Energy Density Solid Cryogenic Rocket Propellants 1: Atoms in Solid  $H_2$ ," PL-TR-93-3014, Phillips Laboratory, Edwards Air Force Base, CA, 1993.

## **Project Objective:**

To identify species produced by HEDM sources

- Background: HEDM species are chemically stable in H<sub>2</sub> matrices in low concentrations and at low temperatures.
- BUT we want higher concentrations (i.e., 5% HEDM in solid H<sub>2</sub>).
- Therefore, knowing the dopant species' identities before and after deposition will result in a better understanding of the deposition process, making it easier to optimize the deposition conditions for high concentration HEDM/solid H<sub>2</sub> matrices.

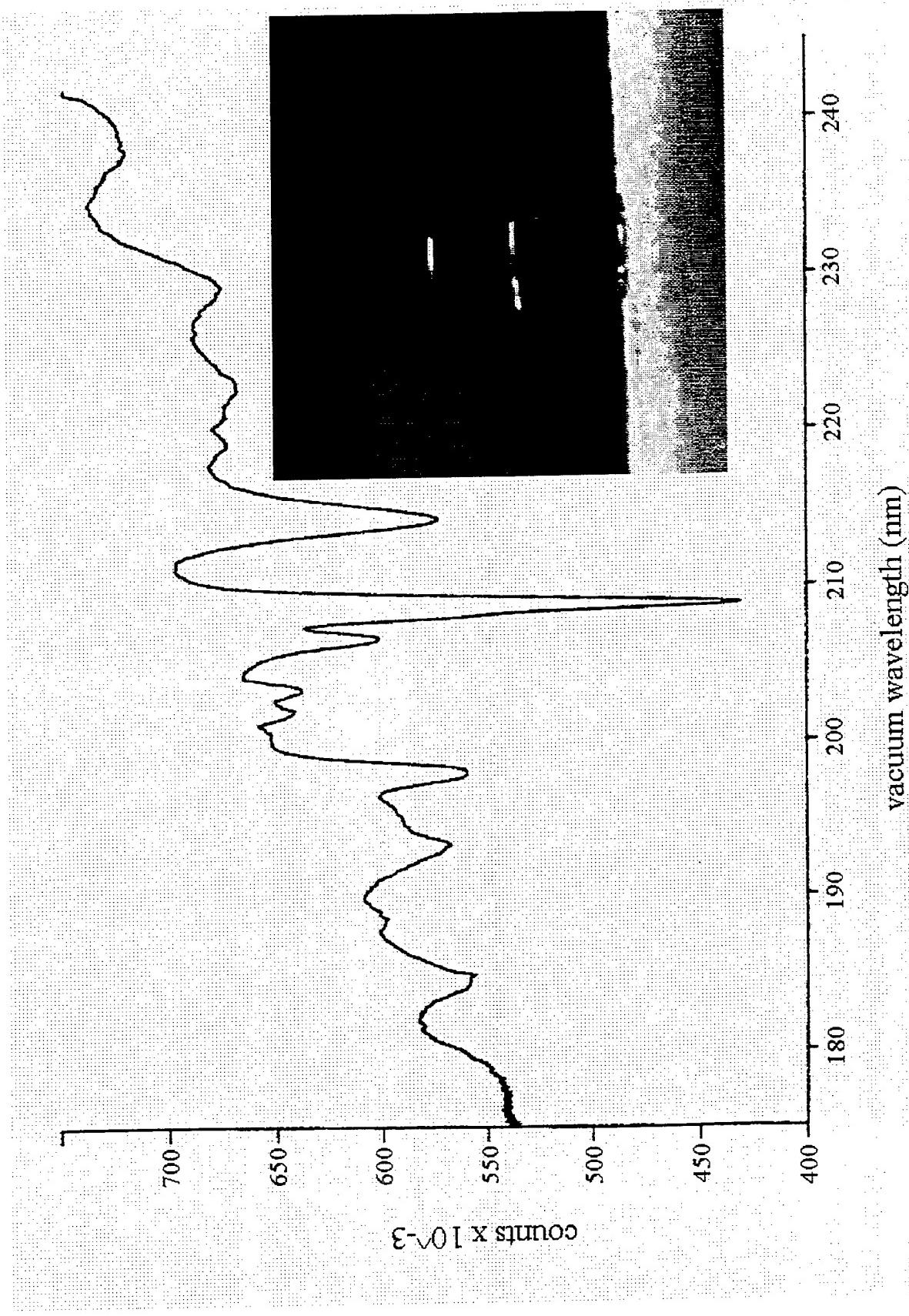
# Sources

## HEDM sources:

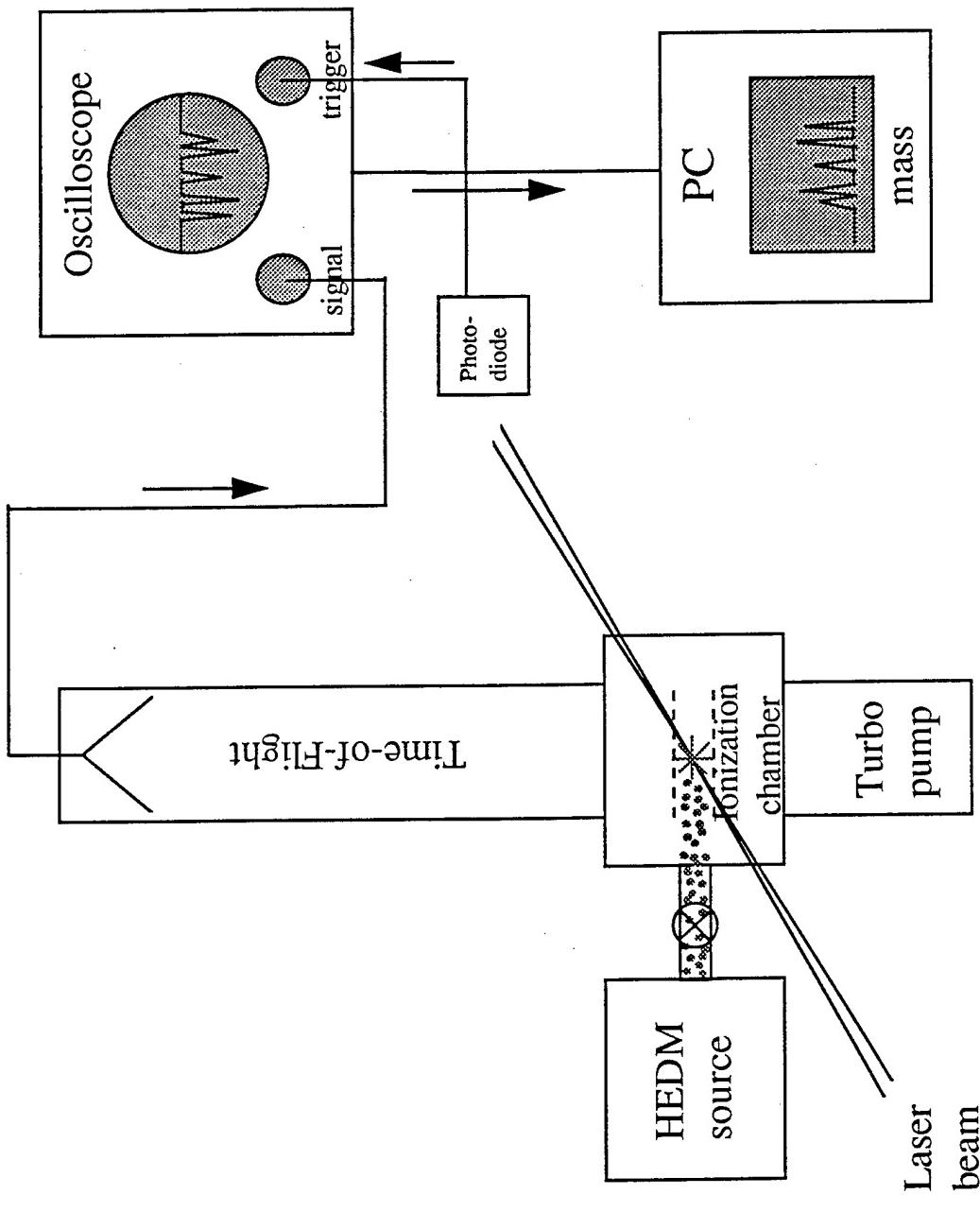
- AFRL-developed boron sources
- commercial sources such as Knudsen ovens or electron beam sources
- tungsten filament sources:
  - made by coating the tungsten filament of a quartz halogen light bulb (Figure 1)
  - advantages: inexpensive, easy and fast to make, suitable for numerous species (metals and solids)

- B sources can easily be made by pyrolyzing  $B_2H_6$  onto a bare filament. It has been proven that B sources made by this technique do produce B atoms.
- However, besides B atoms, other boron-containing species may be produced as well. Or, more generally, any HEDM source may produce many other species besides the target species.

# B atoms in solid argon from B tungsten filament source



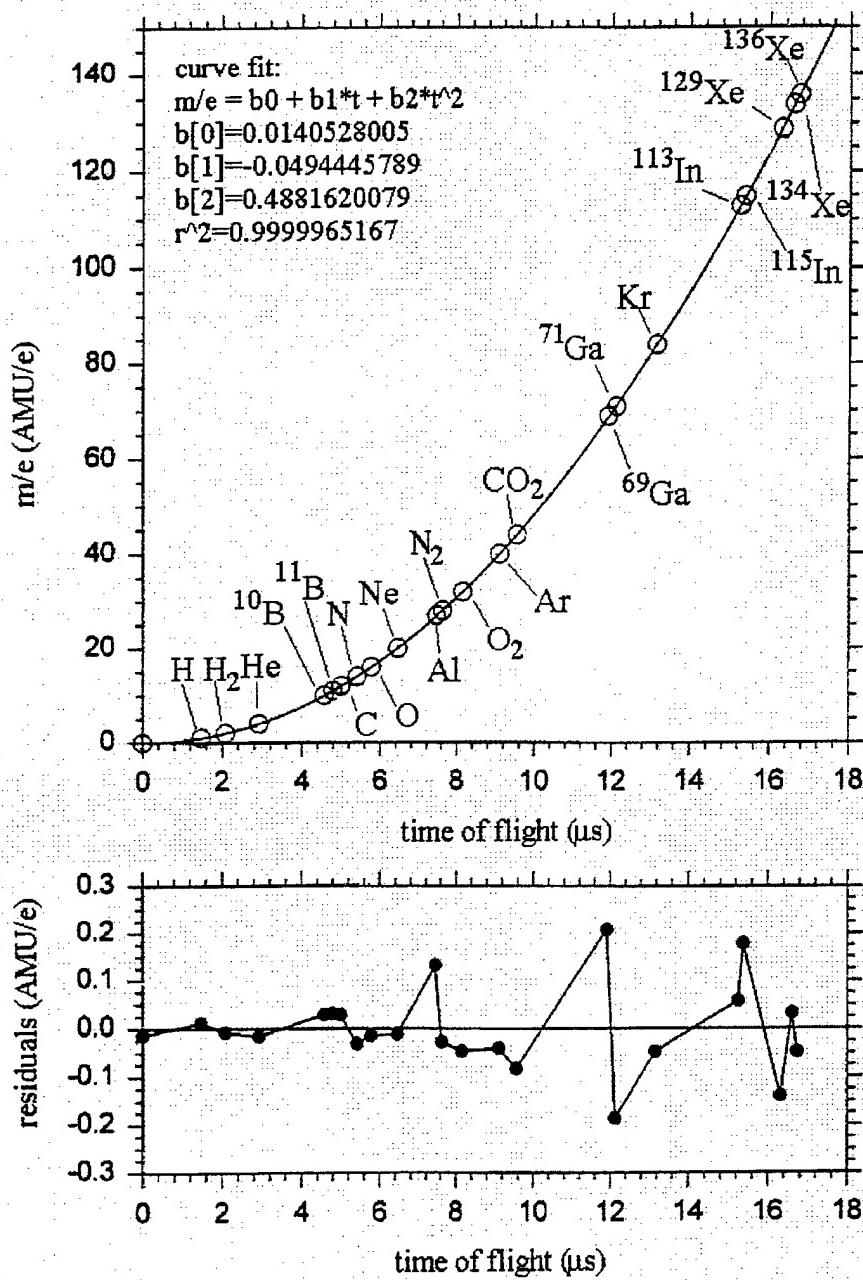
# Time-of-Flight Mass Spectrometry apparatus for HEDM source characterization

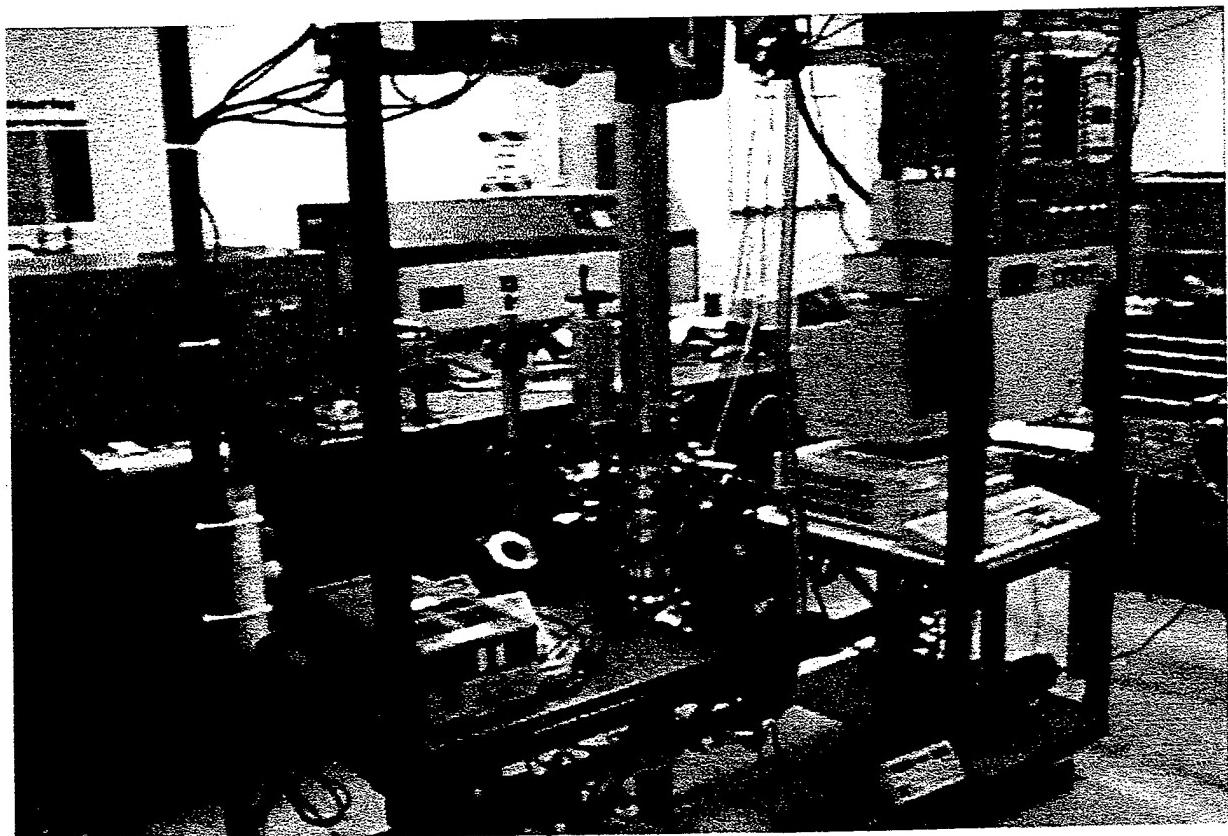


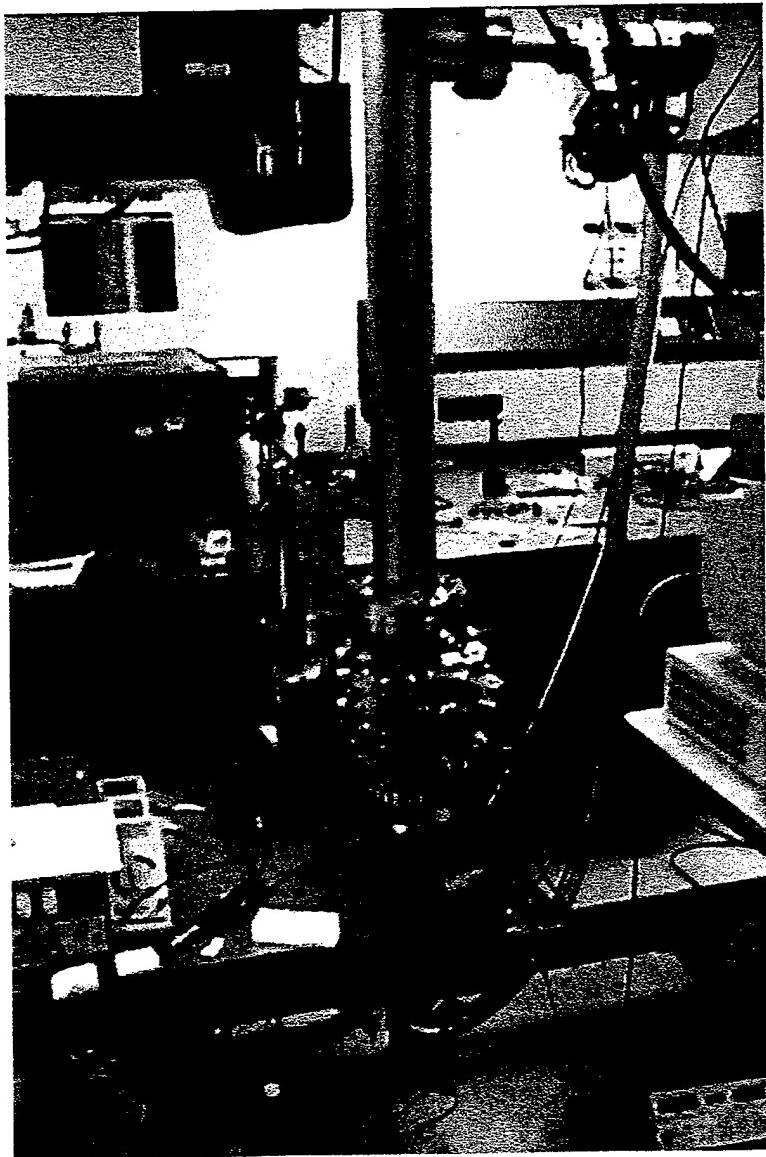
## **Time-of-Flight Mass Spectrometry apparatus for HEDM source characterization (continued)**

1. Source produces beam of HEDM species.
2. HEDM species beam intersects focussed UV laser beam.
3. MPI and MPF take place at intersection → ION SOURCE
4. Ions are accelerated into TOF and detected by microchannel plate detector.
5. Laser scatter onto photodiode gives  $t_0$  → OSCILLOSCOPE TRIGGER
6. TOF signal recorded with oscilloscope, transferred to PC, and converted to mass spectrum.

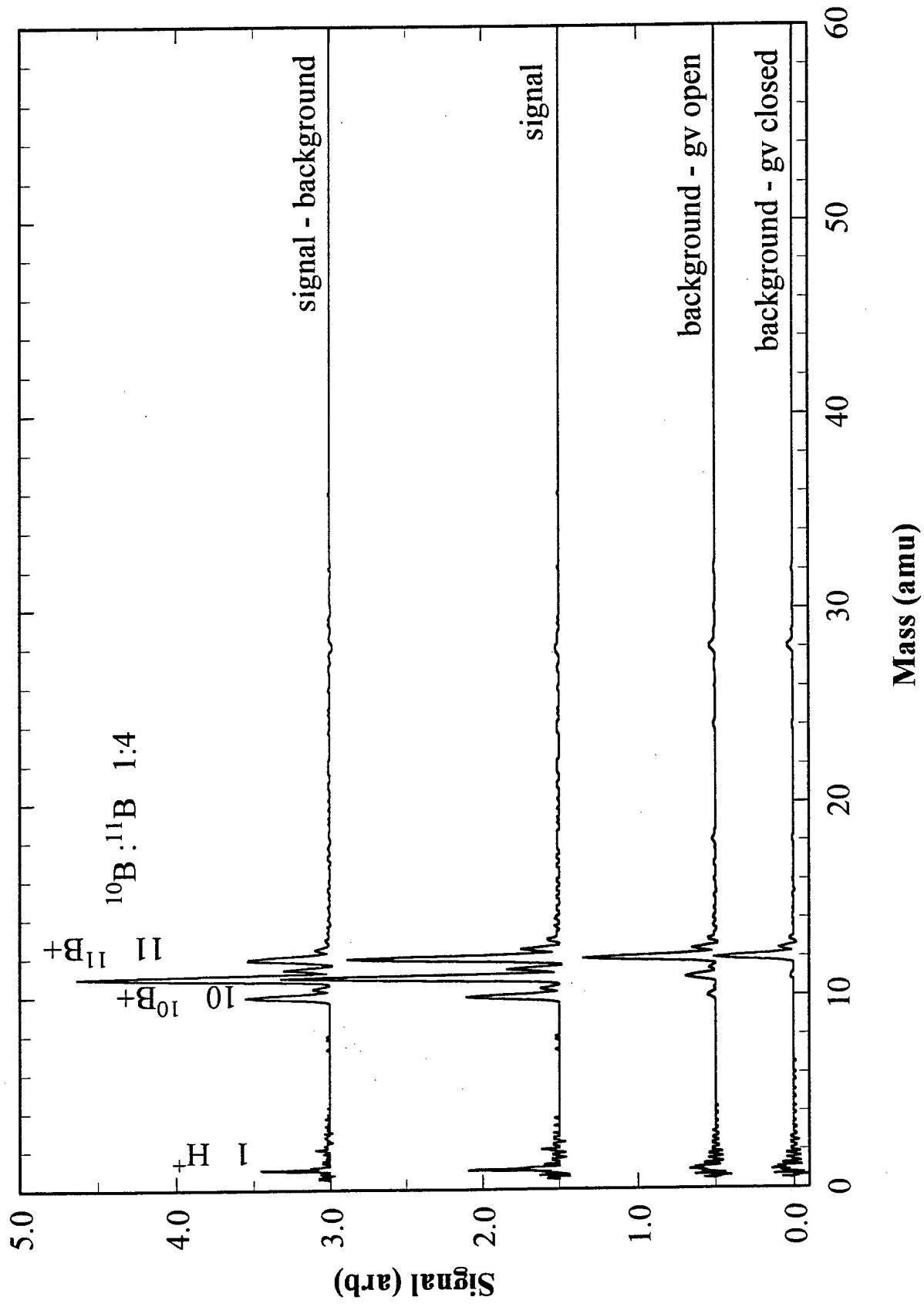
# Ionic mass vs. Time-of-Flight





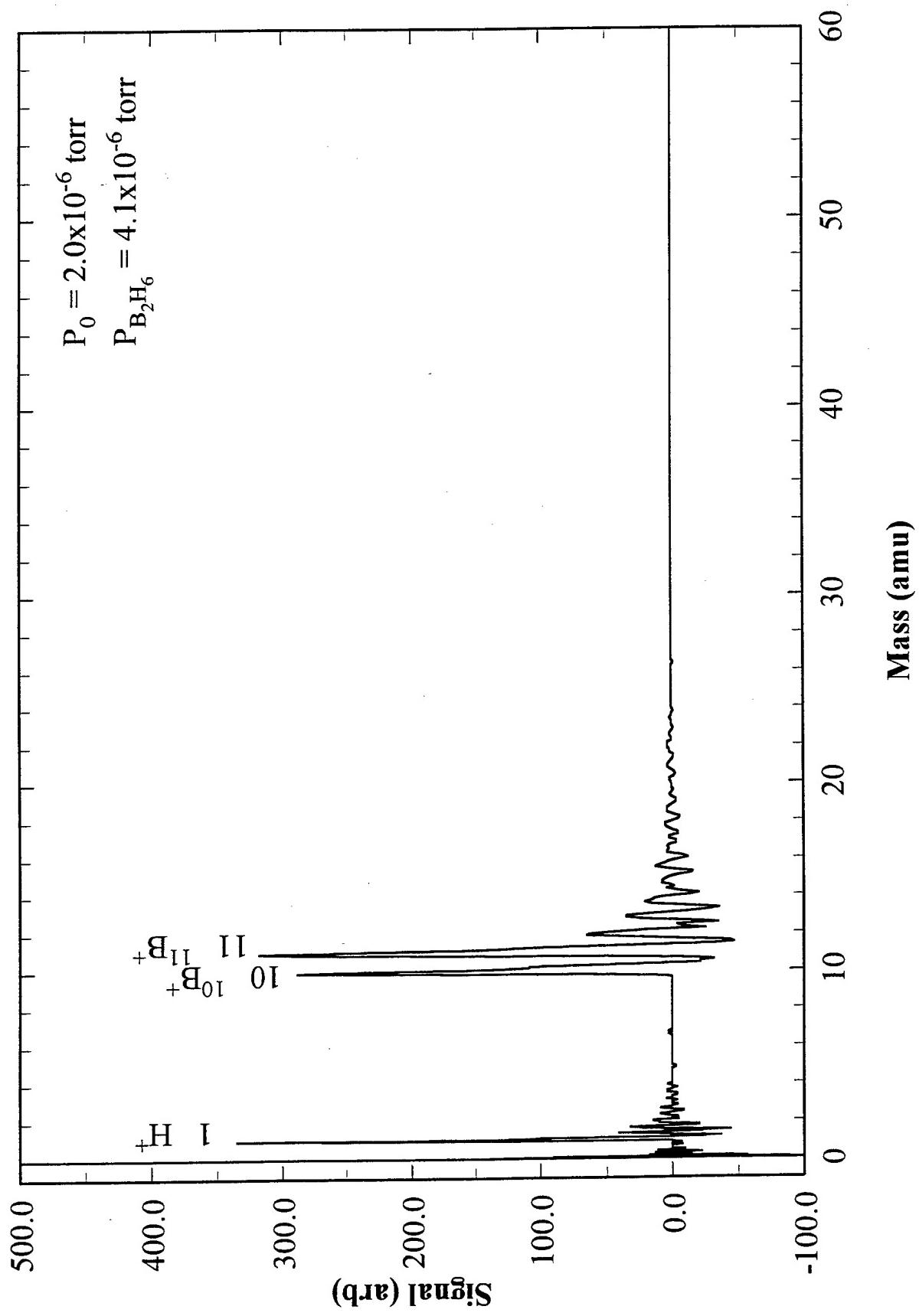


# Boron tungsten filament source, 193 nm



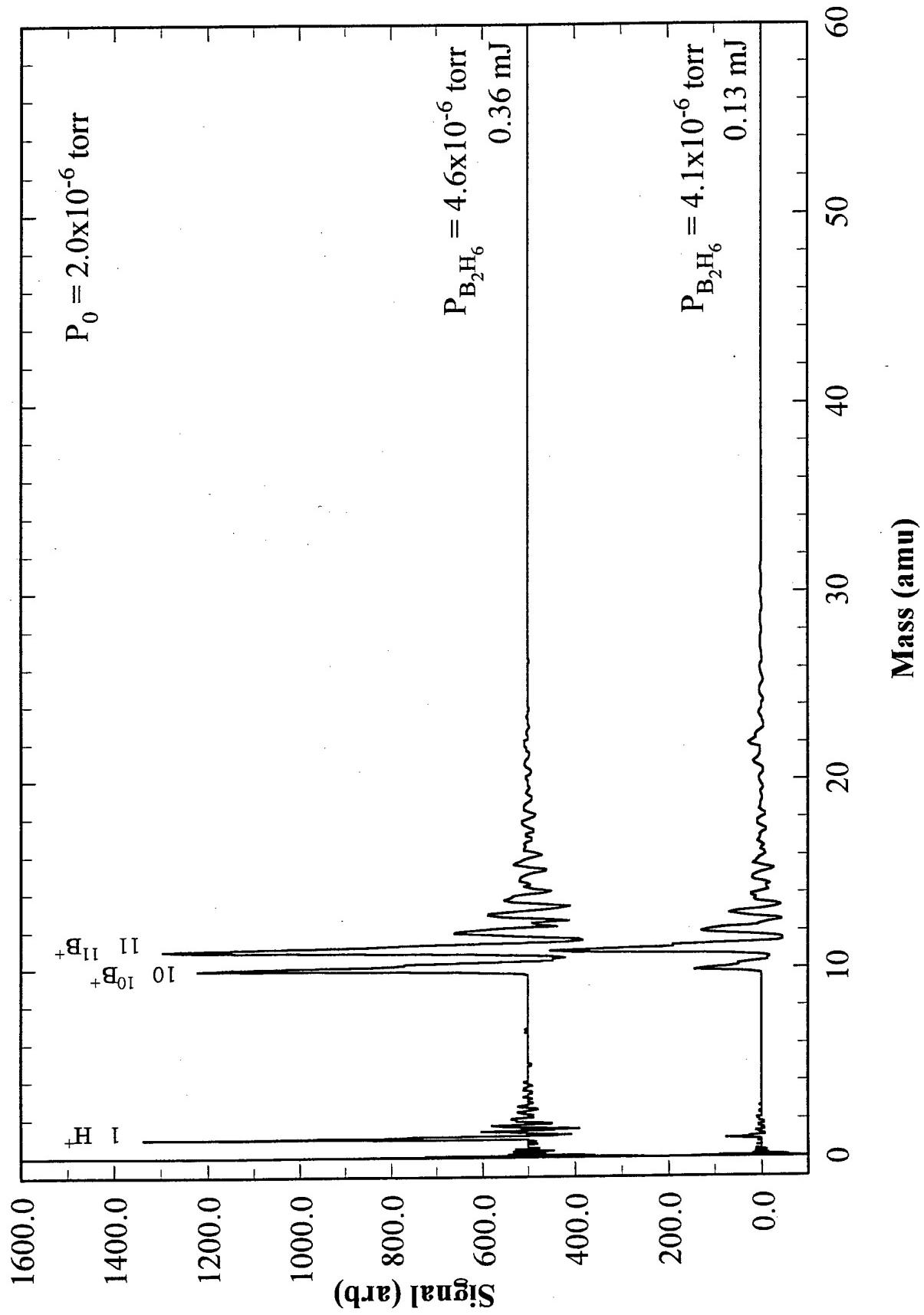
MD213H

# $\text{B}_2\text{H}_6$ , 193 nm



MD230BB

# $\text{B}_2\text{H}_6$ ionized with different intensities of 193 nm light



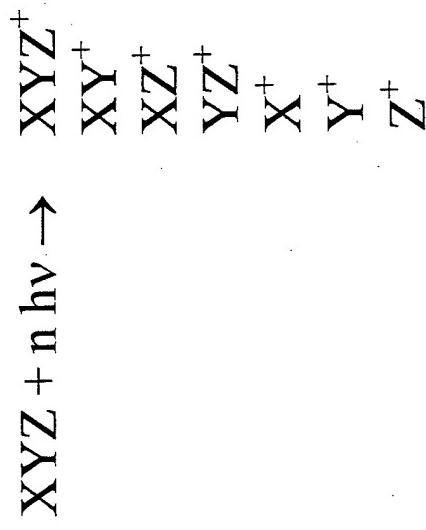
MD230J, MD230BB

# Multi-photon Ionization and Photofragmentation

Ideally for Time-Of-Flight Mass Spectrometry analysis:



In practice:



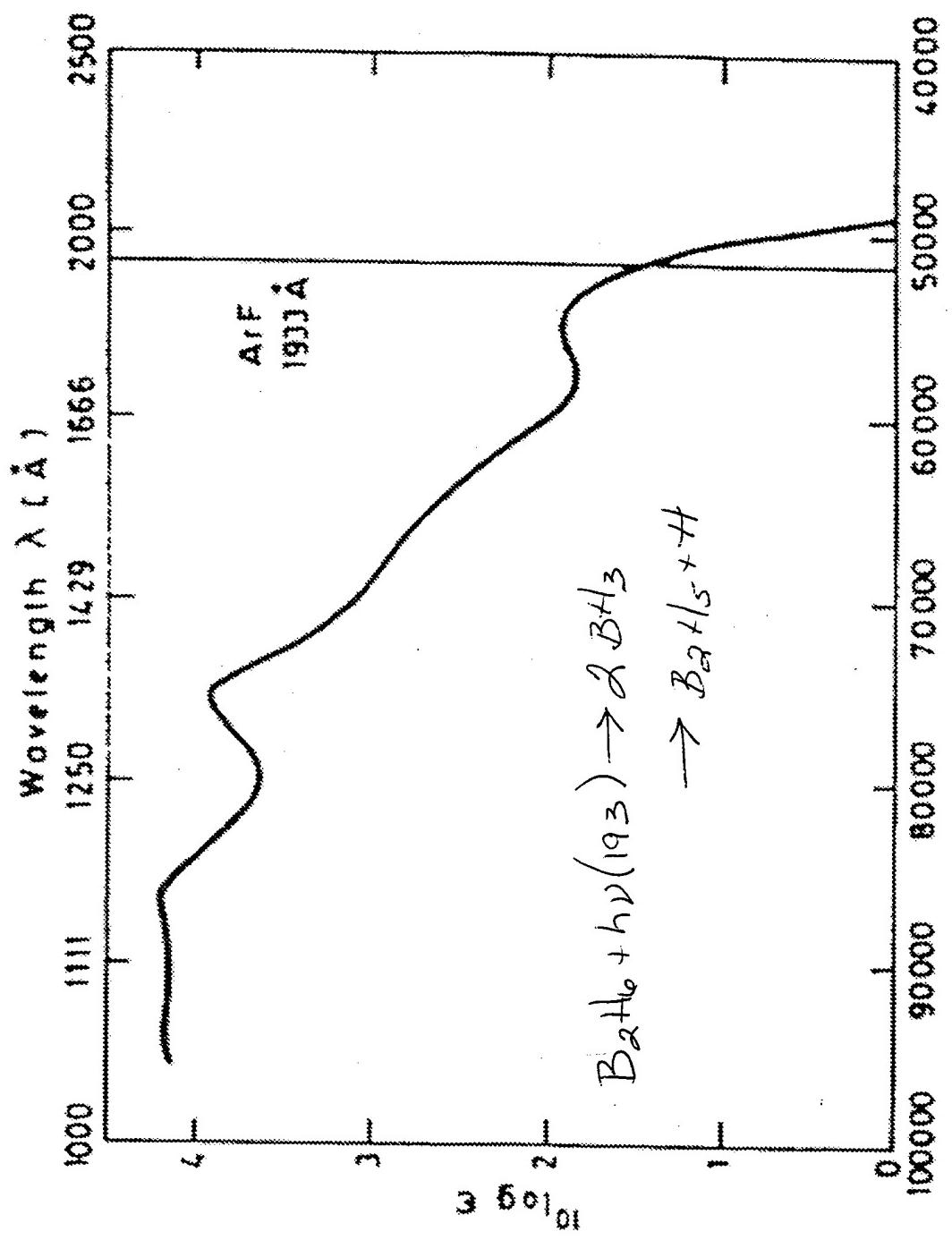
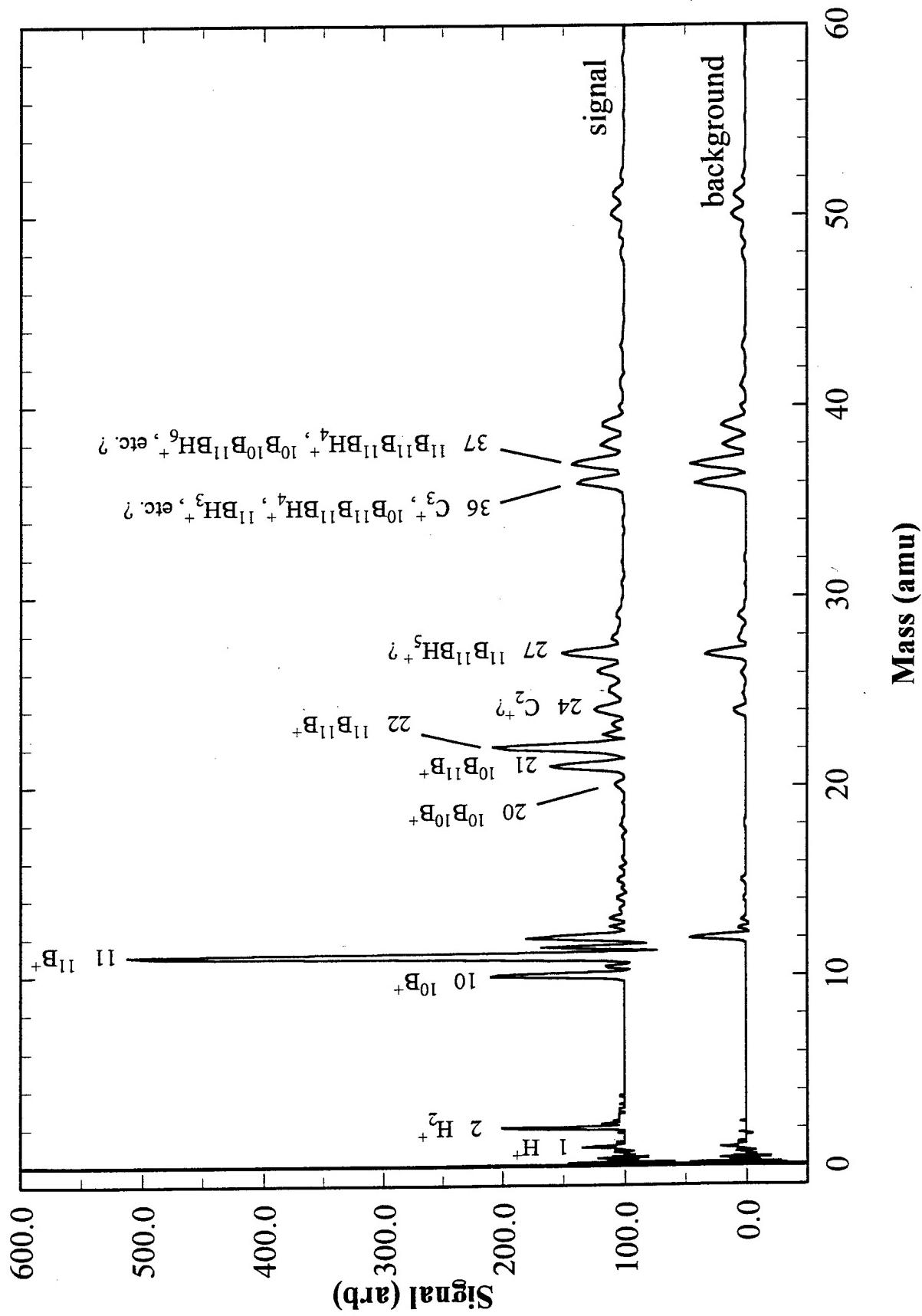


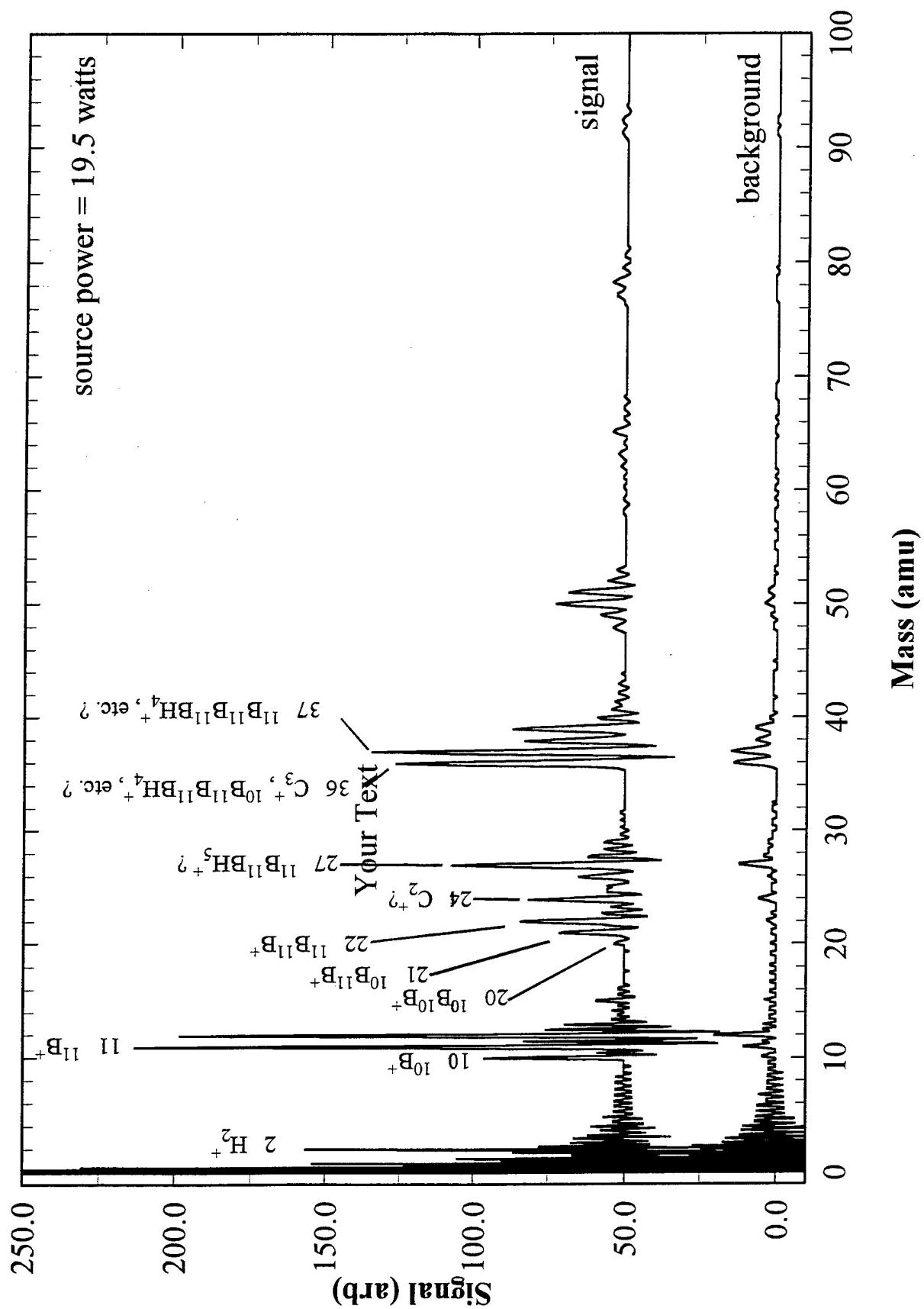
FIG. 1. The UV absorption spectrum of  $B_2H_6$  in the gas phase after W. Fugl<sup>11</sup>

$$\left[ \epsilon = \frac{1}{c \cdot 10} \log \left( \frac{I_0}{I} \right) (1 \text{ mol}^{-1} \text{ cm}^{-1}) \right].$$

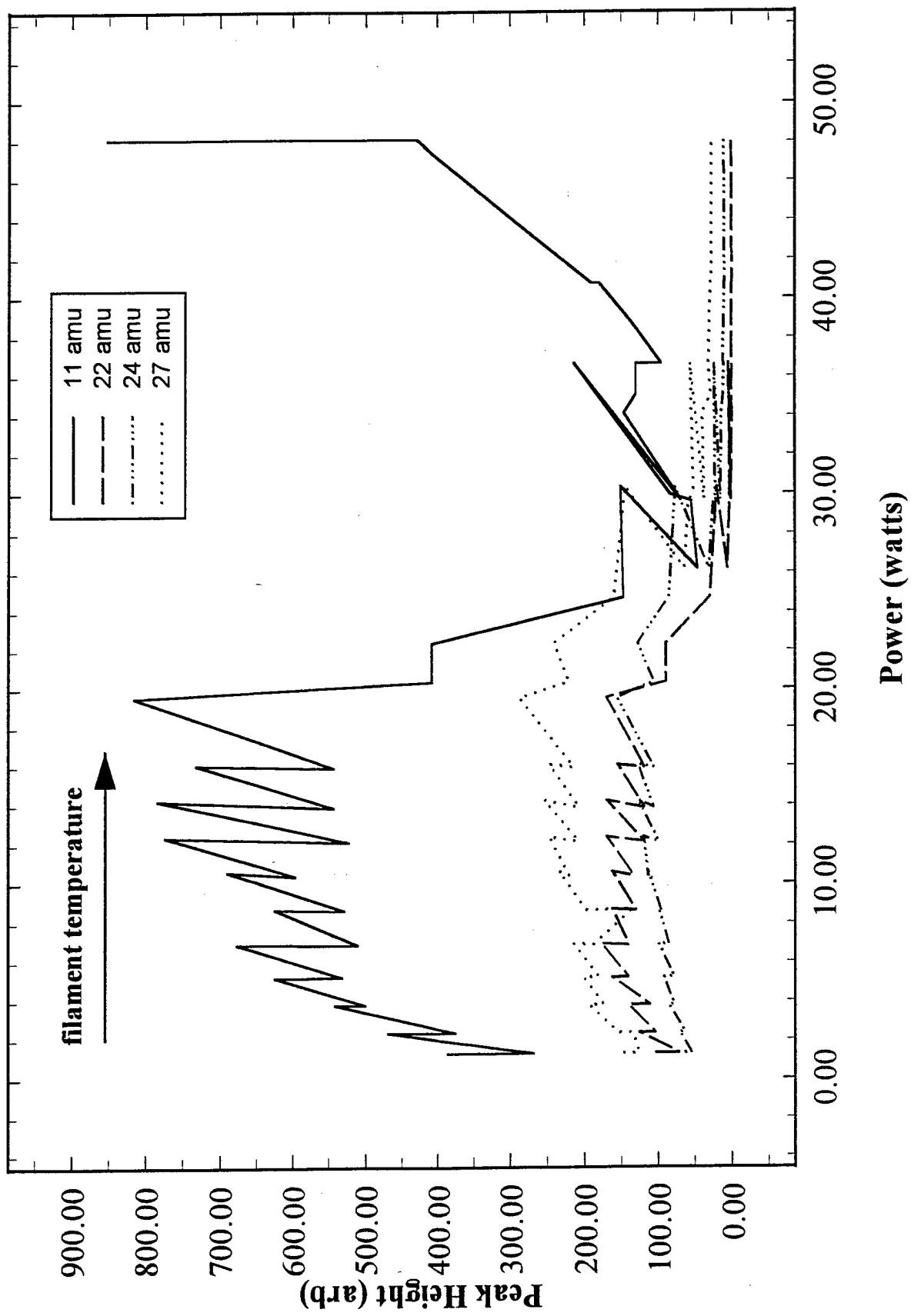
# $B_2H_6$ , 248 nm

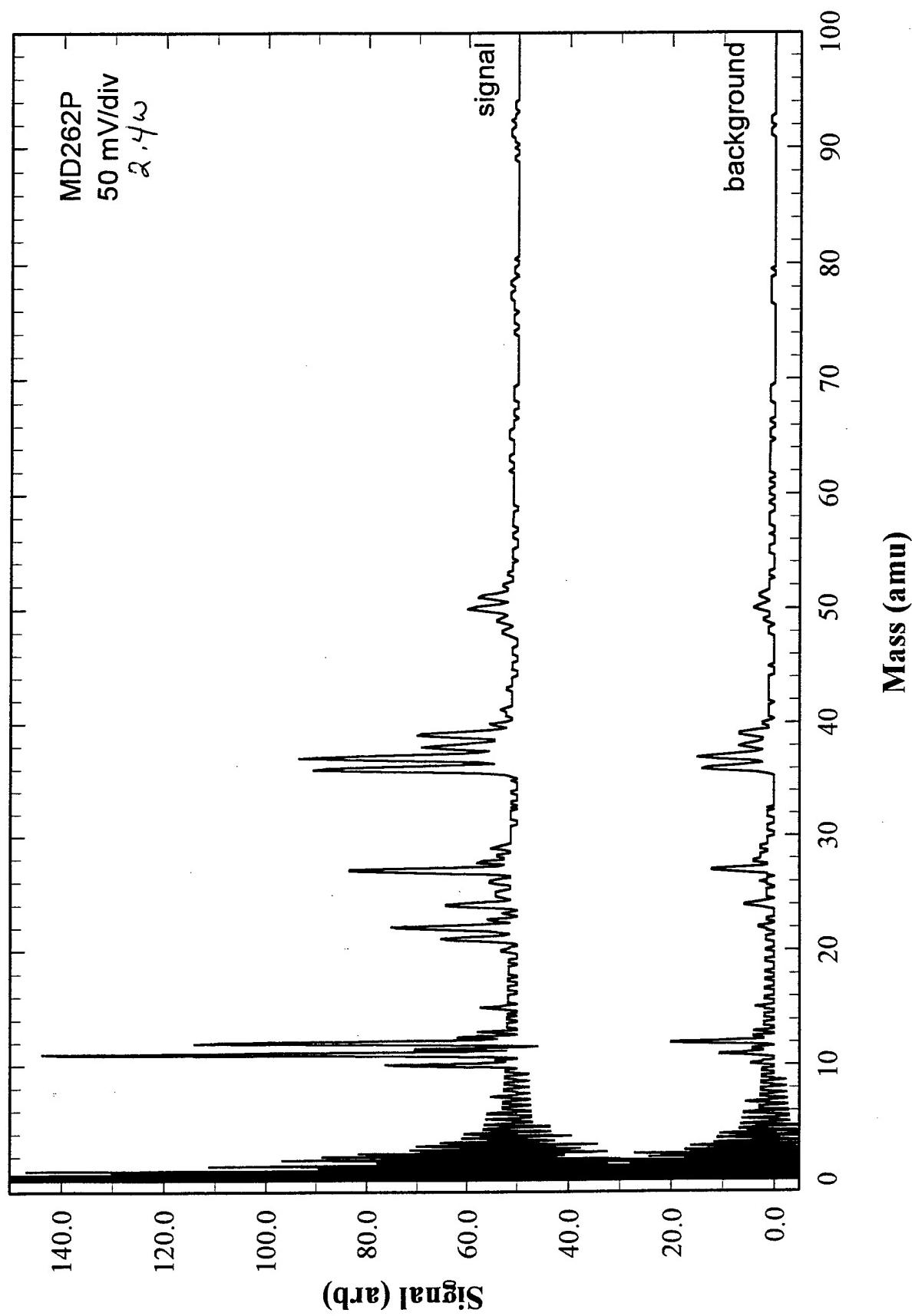


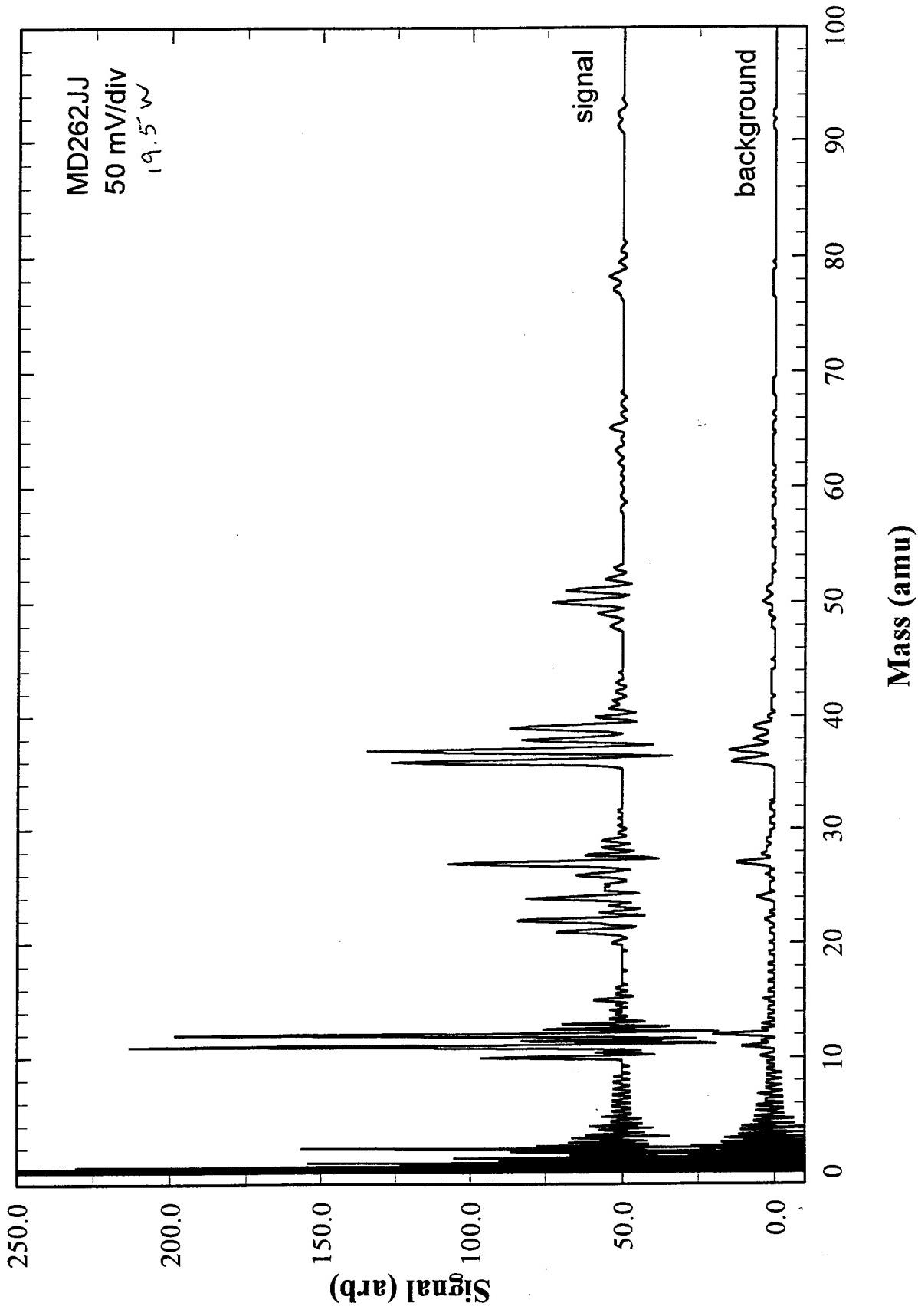
# B source, 248 nm

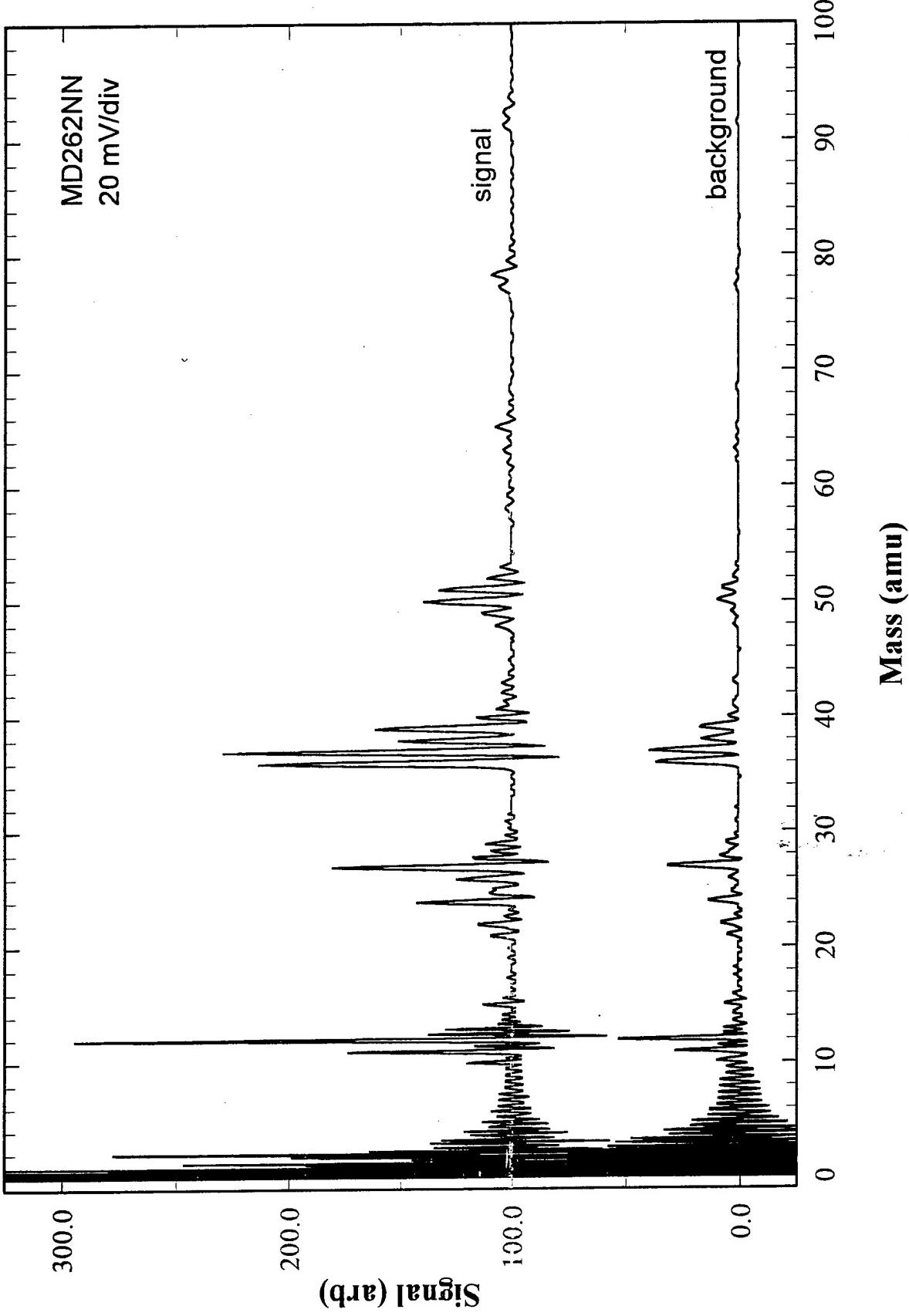


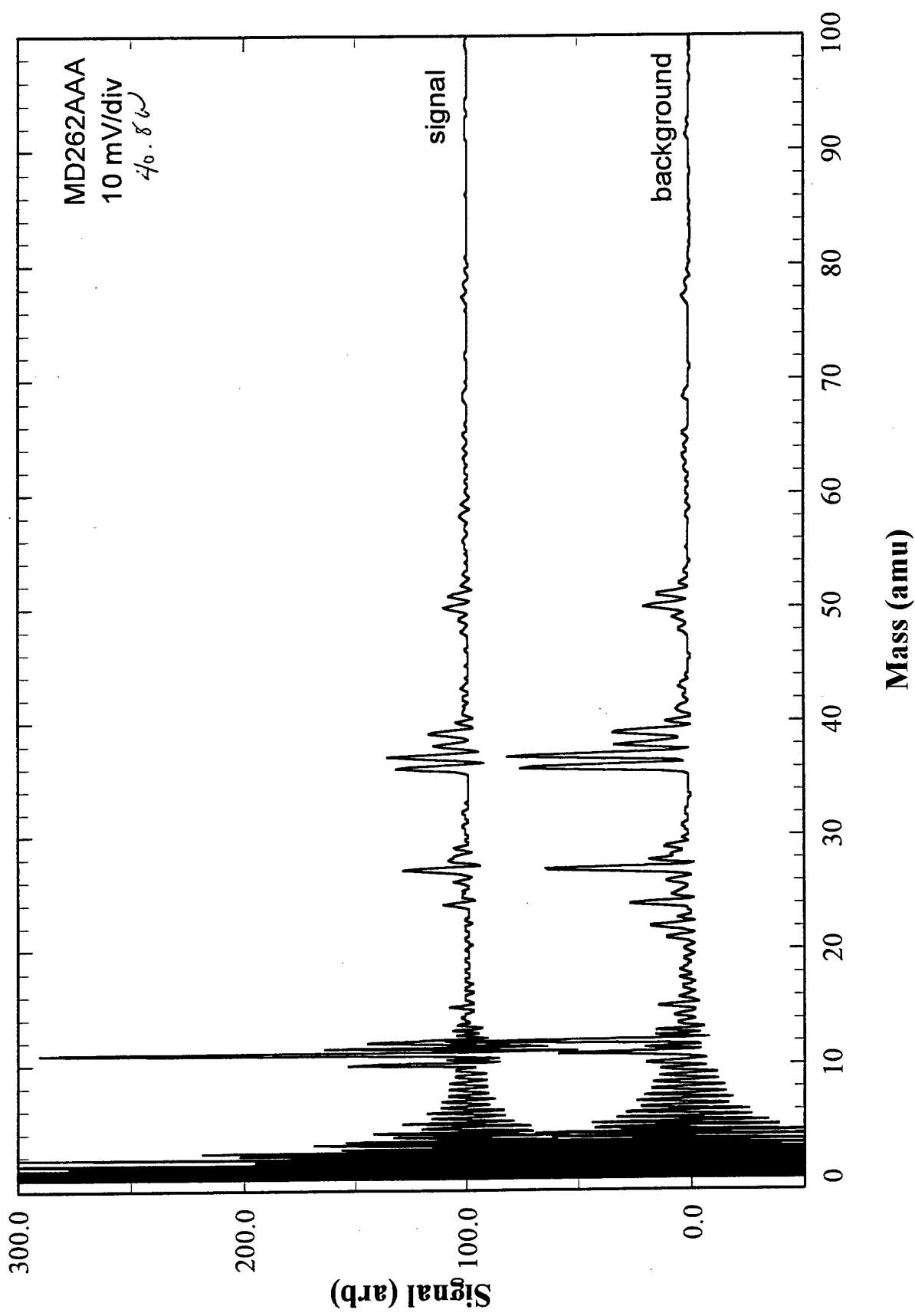
**Peak height vs. Power  
B source, 248 nm**

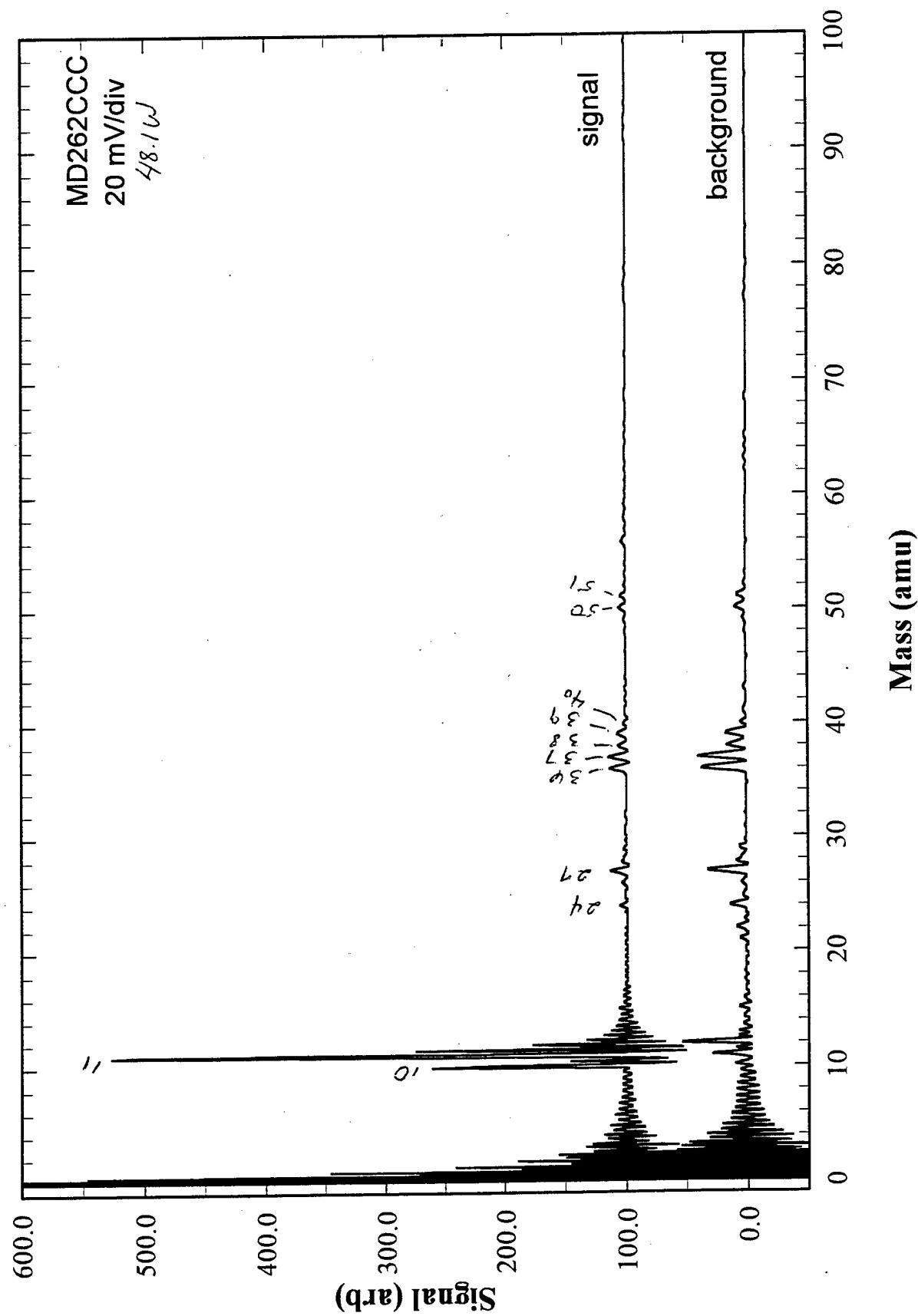












## Summary

- A system to characterize HEDM sources by time-of-flight mass spectrometry has been assembled, calibrated, and tested. Data show that this system is successful in identifying the products of a HEDM dopant source.
- Sources for species such as B, and metals such as Al, In, and Ga can easily be made by coating a tungsten filament.
- Varying the intensity of the laser beam is not a satisfactory way to avoid photofragmentation.
- Varying the wavelength of the laser beam shows much more potential for determining the identity of the parent ion.
- Analysis of boron source data indicates that B atoms detected at low filament temperatures are produced by the photofragmentation of boranes. However, as the filament is heated to higher temperatures, B atoms are emitted directly from the source.

## *(Handwritten)* Future Directions

- Refine detection and ionization scheme:
    - fix electrical ringing in TOF
    - more experiments with different ionization wavelengths
  - Study other sources besides tungsten filament sources
- Ultimate goal:** be able to take a source off the shelf, quickly characterize and/or verify its output, and do a deposition ✓